

#### **Company Presentation**



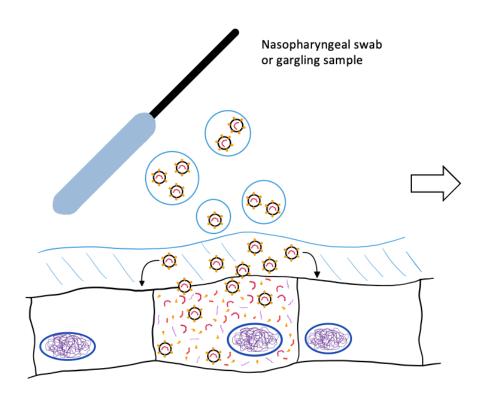
*Marko Poglitsch, PhD Lead Scientific Advisor & Co-Founder* 



**Bernhard Klemen** CEO & Co-Founder

5<sup>th</sup> BioNanoNet Member Welcome Webinar May 25<sup>th</sup> 2023

## **Quarantine and Innovation...**

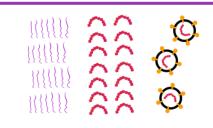


Infected cells of the respiratory epithelium shed virus particles that can infect other individuals via carry over of saliva and aerosol droplets. Samples are obtained via a nasopharyngeal swap or gurgling with saline solution.

#### **Complex samples:**

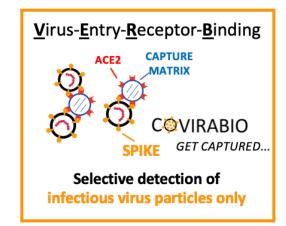
- Epithelial cells
- Cell fragments and mucus
- Virus proteins and fragments
- Free genomic viral RNA
- Intracellular viral mRNAs
- Infectious virus particles

#### **CLASSIC RT-PCR**



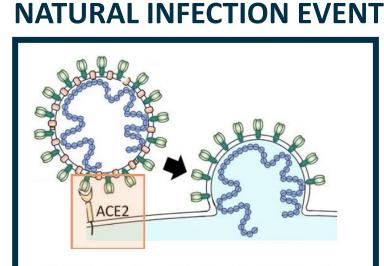
Detection of RNA fragments, free viral RNA and RNA containing virus particles

#### **VERB-ASSAY**

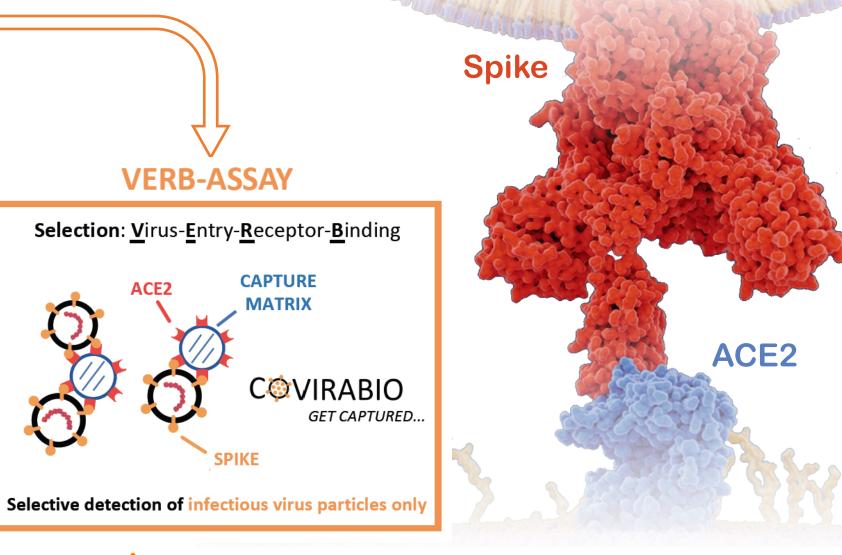




## VERB-Approach: Virus Entry Receptor Binding



**Biochemical background:** Binding of SARS-CoV-2 spike protein to the virus entry receptor ACE2 on airway epithelial cells is the key step in SARS-CoV-2 infection.



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## VERB-Approach: Proof-of-Concept Model SARS-CoV-2

**Spike** 

ACE2

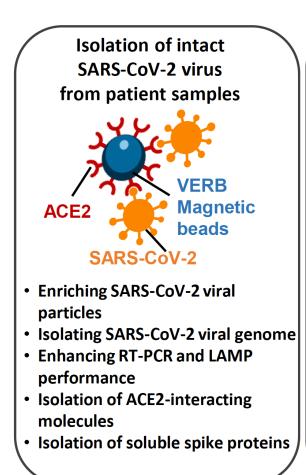
• Detection of

#### **INTACT VIRUS PARTICLES ONLY**

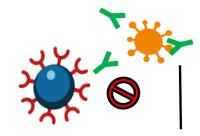
- Sample material: swabs, body fluids
- Multiple detection options
  - + Colorimetric / Fluorimetric
  - + (*RT-q*)*PCR*
  - + SPR (Surface-Plasmon-Resonance)
  - + Biosensors
  - + Microfluidics
- Completed p.o.c. in COVID-19 (PFU vs. VERB)

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## SARS-CoV-2 VERB Kit



Detect the presence of SARS-CoV-2 neutralizing antibodies



#### **Neutralizing antibodies**

- Determining viral neutralizing titers
- Immune response to variants of concern
- Vaccine efficiency monitoring
- Screening for SARS-CoV-2 viral entry inhibitors

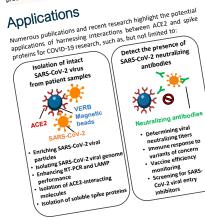


Virus Entry Receptor Binding – VERB assay (RUO) A simple tool to isolate intact SARS-CoV-2 particles and detect the presence of neutralizing antibodies Procedures

#### Assay Principle

The highly efficient binding of SARS-COV-2 via its spike protein to the highly encient binding of SARS-CUY-2 via its spike protein of its cellular entry receptor ACE2 is the basis for the successful initiation of the infection cycle of this virus and forms the initiation of the infection cycle of this virus and rounds are molecular principle of the VERB (Virus Entry Receptor Binding) noneural principle of the vero (virus tiny receptor oniums) approach developed by Covirabio. A capture matrix has been approach aeveroped by Covirable. A capture match has been developed which allows the highly efficient isolation of intact SARS-CoV-2 particles and the determination of the presence of neutralizing antibodies in a routine lab environment. The VERB assay is a stand-alone sample preparation method

the veno assay is a static-aroute satisfic preparation include compatible with downstream detection methods such as RT-qPCR winipauvie with womistream verection methods such as Ri-4PCR and thus can be easily integrated into existing laboratory procedures.



Key Features

Free of viral RNA fragments and deb

Magnetic beads functiona

Order information Cat. COV007-RUO

ant human ACE2

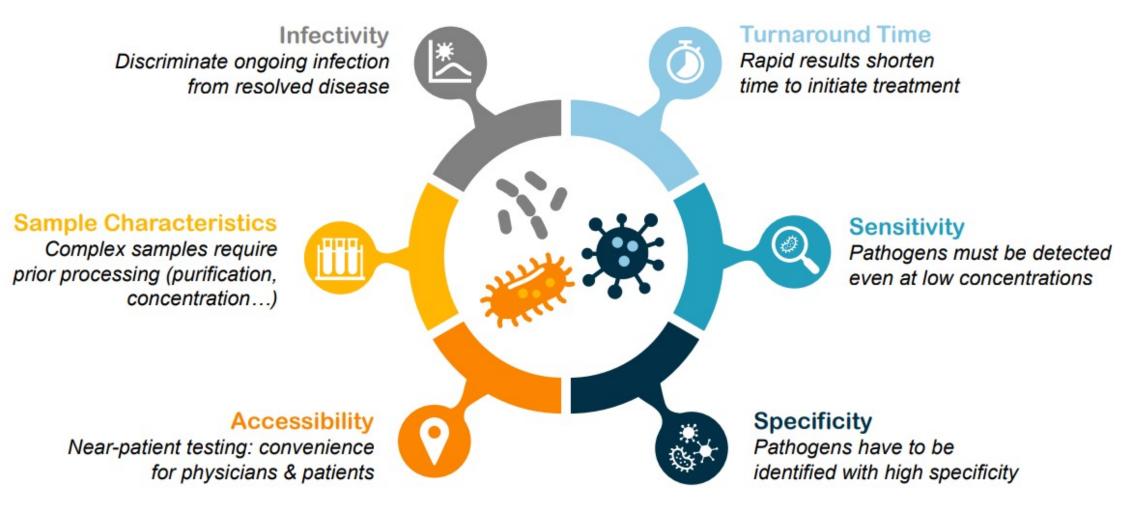
#### Viral capture ≤ 30 min Viral RNA extracted Automatic mode (e.g. Maelstrom and King Fisher systems) for medium and high throughput Manual mode for low sample numbers Performance 25-20-Fig. 1 Data supporting the isolation of intact virus particles rig. 1 uata supporting the isolation of interview particles with VERB beads. Heat inactivation of a SARS-CoV-2 containing build be the state of the WITH VERIS DEBUGS. Heat inactivation of a SARS-COV-2 containing Clinical sample abolishes the isolation of RNA captured by VERB clinical sample abousties the isolation of NNA captured by Verb beads (< 5% input), when compared to the captured RNA from tesaus (< 3% input), when compared to the captured KNA from the sample kept on ice (ca. 35% input). The data is normalized the sample kept on ice (ca. 35% input). The uata is normalized to the total RNA of the sample without VERB bead capture process (input). Capture and enrich only intact virus

Add VERB beads

Nasal swab, Serum

Fig. 2 Presence of neutralizing antibodies can be detected ris, a reserve or neuranens announces can be deterring with the VERB beads. Reference serum with neutralizing will the verb beaus, neterence serunt with reutaniants antibodies against SARS-CoV-2 (EURMO17) was serially diluted and incubated with SARS-COV-2 pseudovirus before unueu anu muualeu wan sanouve peuvernus euror subjection to the VERB assay. The titer of neutralizing anti-discuss determined between bask control by VERB subjection to the VERB assay. The titer of neutralizing antibodies was determined how much RNA captured by VERB beads. The IC<sub>50</sub> determined by VERB was at 47.3. COVIRABIO Further information info@covirabio.com

## **Challenges in the Diagnosis of Infectious Diseases**

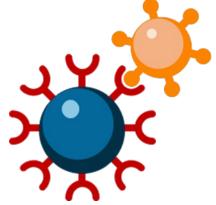


Clock

## **MISSION**

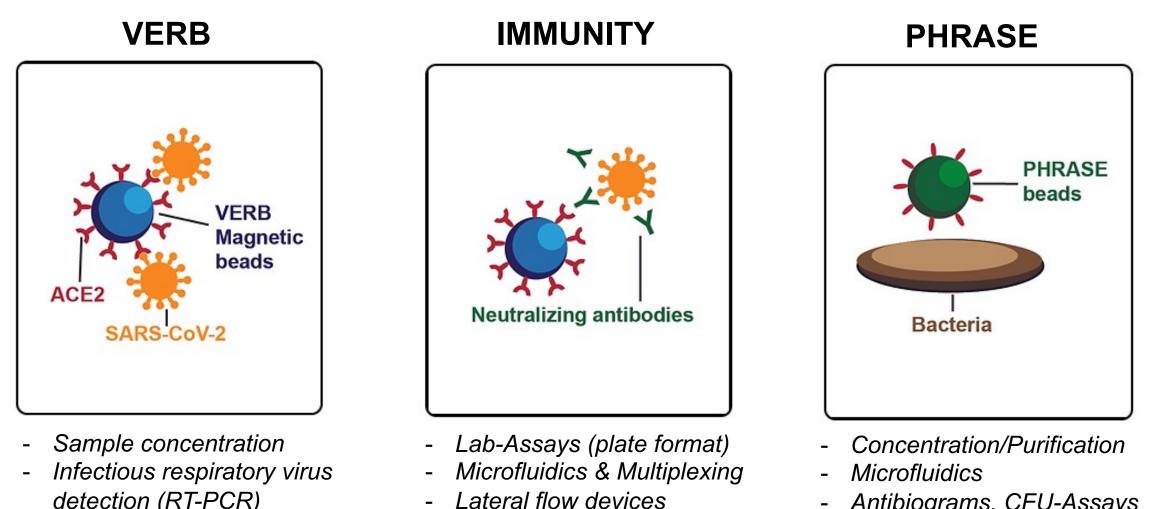
## **Diagnostic Utilization of Host-Pathogen Interactions**

- identify **specific and immune-relevant** host/pathogen interactions
- develop novel highly sensitive nanoparticle-based diagnostic tools
- implement emerging technologies to develop improved and affordable diagnostic solutions for pathogen detection and immunity monitoring (microfluidics, biosensors,...)
- providing robust and sensitive pre-analytic approaches for pathogen detection ("enrichment")
- offer tools for rapid and sensitive detection of infectious respiratory viruses





## **Platform Technology: Functionalized Nanoparticles**

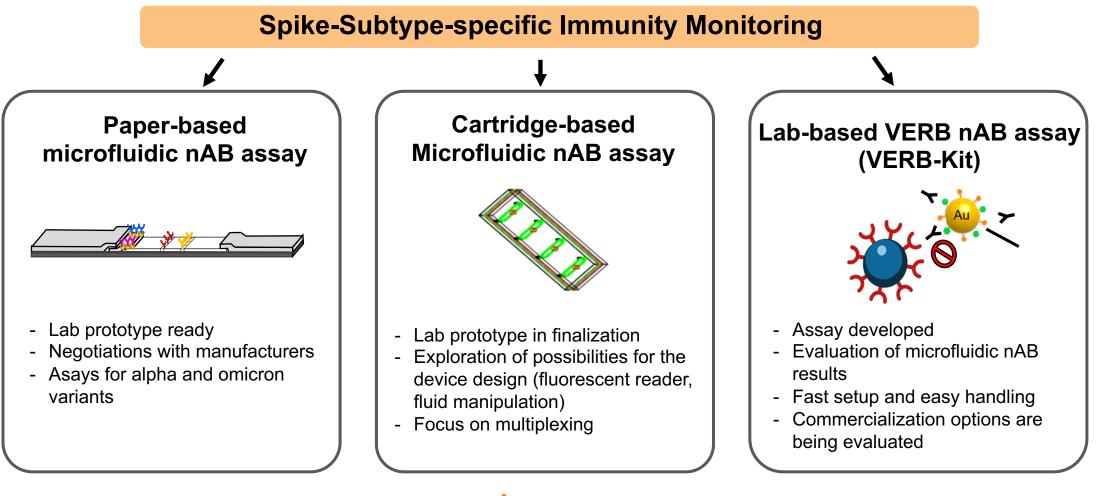


RABIO

detection (RT-PCR)

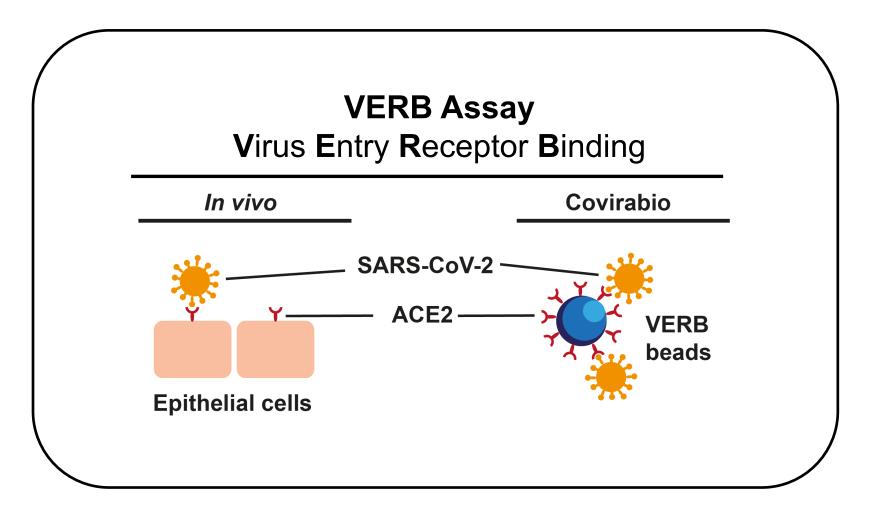
Antibiograms, CFU-Assays

# *Immunity Monitoring:* SARS-CoV-2 as a p.o.c. Model Nanoparticle based Detection of Neutralizing Antibodies



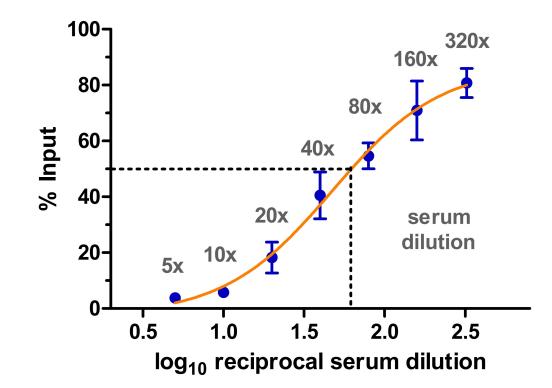
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## SARS-CoV-2 Immunity Monitoring





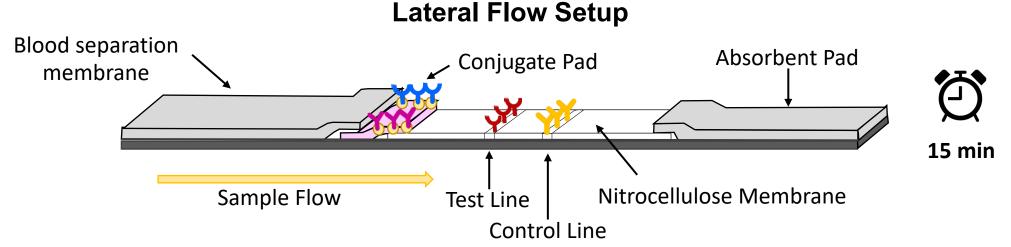
## SARS-CoV-2 Immunity Monitoring (VERB-Kit)



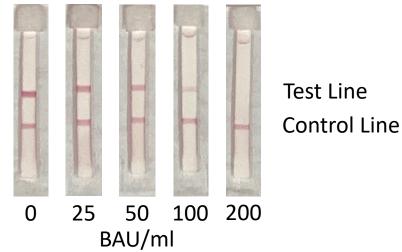
**Presence of neutralizing antibodies can be detected with the VERB beads.** Reference serum with neutralizing antibodies against SARS-CoV-2 (EURM017) was serially diluted and incubated with SARS-CoV-2 pseudovirus before subjection to the VERB assay. The titer of neutralizing antibodies was determined by how much VERB-captured RNA could be detected. The IC<sub>50</sub> determined by the VERB kit was at serum dilution of 47.3.



## *Immunity Monitoring* SARS-CoV-2 Nanoparticle-based Detection of Neutralizing Antibodies



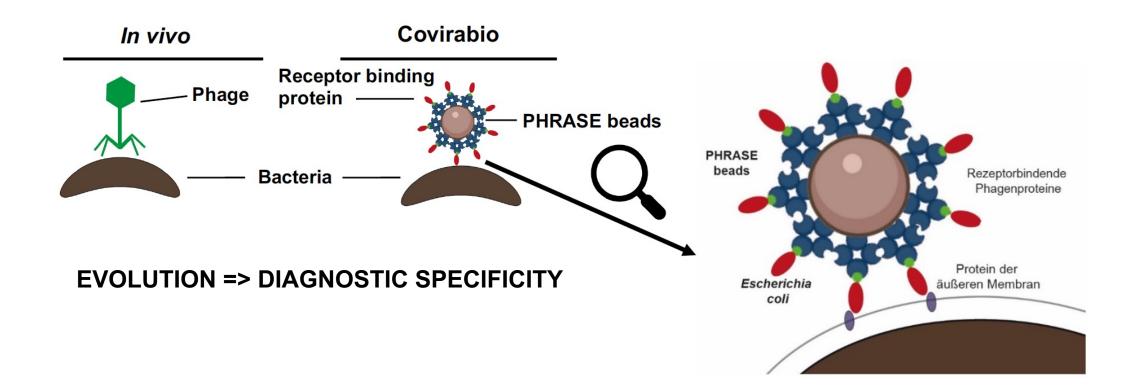
#### **Functional Lab-Prototype**



- Rapid test for immunization status against SARS-CoV-2
- Optional analysis of whole blood or serum
- Based on interaction of **natural receptor/ligand pairs**
- BAU sensitivity optimized for self-testing

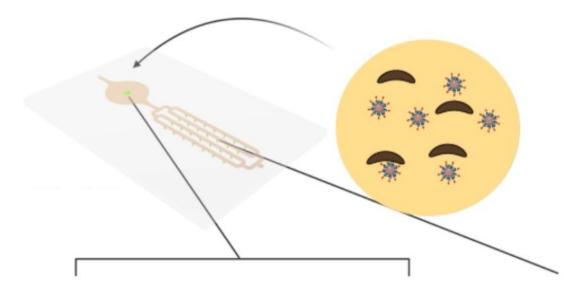
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## **PHRASE** Approach <u>PH</u>age-<u>R</u>eceptor-<u>A</u>ssisted-<u>S</u>ample-<u>E</u>nrichment

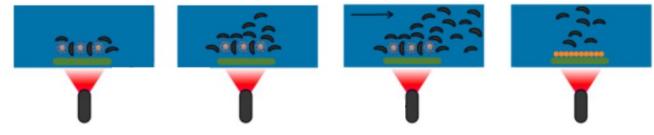




## **PHRASE Microfluidics**



In-cartridge cultivation and growth monitoring



Microfluidic-based real-time detection of pathogenic bacteria in biological samples

=> Qualitative Diagnostics=> CFU-Determination=> Antibiograms



## **Team and Partners**

#### R&D Team



#### Scientific Advisory Board Members



#### Prof. Dr. Peter Ertl

- Biotechnology; Chemistry (PhD); biophysicist (postdoctoral);
- Technical University of Vienna; Berkeley (Fulbright Scholar);
- Research focus on biosensors, microfluid devices, lab-/ organ-on-a-chip technologies

#### R&D Partners





Wiener Gesundheitsverbund Für die Klinik Donaustadt



#### **Private-Sector Partners**



 Diagnostics company with 10+ year ACE2/ analytics experience
Lab environment/ staff